

# PATENT ABSTRACTS OF JAPAN

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(71)Applicant: AGFA GEVAERT AG

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(72)Inventor: WERNICKE UBBO

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# (54) RAPID DEVELOPING METHOD OF COLOR MATERIAL

#### (57)Abstract:

PURPOSE: To enable rapid development of a silver halide recording material by treating the photographic material with a developer or a soln. I of its salt and then subjecting the photographic material impregnated with the developer to the treatment with an alkali-contg. soln. II.

CONSTITUTION: The photographic material has photosensitive silver halide emulsion layers having at least three different spectral photosensitivities on a base body, and the cyan coupler, magenta coupler and yellow coupler in the layers are spectrally continuous. The recording material is exposed for an image and treated with a developer or its salt soln. I under the condition of pH1 to 8 for 1 to 10sec. The photographic material impregnated with the developer is treated with an alkali-contg. soln. II under the condition of pH10 to 14 for 1 to 10sec. for development. Thereby, partial images of cyan, magenta and yellow are formed and combined to give a final color image. Thus, rapid development of a color photographic material is realized.

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❷発明の名称

カラー材料の迅速現像方法

**劉特 顧 平1-70788** 

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優先権主張

70発 明 者

ウポ・ベルニツケ

ドイツ連邦共和国デー - 5000ケルン91・ラターマウスプフ

アート 21

勿出 願 人 アグファーゲヴェル

ドイツ連邦共和国レーフエルクーゼン(番地なし)

ト・アクチエンゲゼル シヤフト

四代 理 人

弁理士 小田島 平吉

#### 朔 解 者

1. [発明の名称]

カラー材料の迅速現象方法

2. [特許請求の範囲]

1. シアンカブラー、マゼンタカブラー及び黄カブラーがそれぞれスペクトル的に適合されている 少くとも三つの異なるスペクトル感光性を有する 感光性ハロゲン化級乳剤層を層支持体上に含有す る固像的に露光されたハロゲン化級記機材料を次 の処理段階:

- (1) 写真材料の現象剤またはその塩の静液 I による、p H 1 ~ 8 における 1 ~ 1 0 秒間の処理、
  - (2) 現像剤で合設された写真材料のアルカリー合有溶液 I による、p H 1 0 ~ 1 4 における 1 ~ 1 0 秒間の処理、

に付することを特徴とするハロゲン化銀配録材料 の迅速現象方法。

3. [発明の詳細な説明]

本発明は塩化銀高合量の写真材料の迅速現像に特に渡した方法に関する。

本発明を要約すれば、シアンカプラー、マゼンタカプラー及び黄カプラーがそれぞれスペクトル的に連合されている少くとも三つの異なるスペクトル感光性を有する感光性ハロゲン化級乳剤層を層支持体上に含有する、画像的に露光されたハロゲン化級記録材料を次の処理段階:

- (1) 写真材料の現象剤またはその塩の溶液 1 による、p H 1~8における1~10秒間の処理、
- (2) 現象剤で含張された写真材料のアルカリー含有溶液 ■による、pHIO~14における1~10秒間の処理、

に付することから成る迅速現像方法であって、これはさもなければ 4 5 秒間以上の常用の現像によってのみ得られる種類の最大カラー密度を生成する。

現象剤酸化生成物とカプラー分子との関像的カプリングによるカラー写真関像の形成はよく知られている。この方法において、シアン、マゼンタ及び費の部分関像が通常生成され、それが組合わさって最終カラー関像が形成される。現像剤は一般に一級芳香族アミノ化合 でありこれは変換さ

(2)

れて現像剤酸化生成物を露光されたハロゲン化級 の個域中に難感する。

用いうるハロゲン化級は臭化級、臭化沃化級、塩化銀、塩化臭化級及び塩化臭化痰化酸酸糖料ない。塩化物高含量のハロゲン化凝乳剤または純粋な大性の過程を表現である。では、その理由はそのようなないのである。即ちそれらの可視光線に対する低アイルの高含量の乳剤は多くの可視光線に対する低アイルのである。即ちそれがアイルなまたはなのである。のは、例えばネガアイルなくことので、毎日のは、例えばネガアイルを製造するのに、増加を表現である。である。塩化物イオンは臭化物イオンの如き抑制的効果を有しないからである。

塩化物含量>95モル%のハロゲン化値乳剤を 臭化物のないカラー現像器と組合せて使用するこ とによりカラー紙の現像時間を210秒から45

2. 現象剤の譲度が上方にある層による消費に 基づき十分に高くない。

3. 層重への現像剤と同時に到達するところの 層I及びまからのハライドの多量の存在はそれに 相応して現象工程に抑制効果を有する。

これらの不利は極度に短かい現像時間(例えば <20分)による迅速角理操作において特に認め られそして常弦による現像を、不可能ではないに せよ、一層困難ならしめる。

今回、一つの特種の現象方法が上記不利を克服するばかりではなく実用上における付加的利点を も与えることが見出された。

本類明は、シアンカプラー、マゼンタカプラー 及び黄カプラーがそれぞれスペクトル的に適合さ れている少くとも三つの異なるスペクトル感光性 を有する感光性ハロゲン化模丸利用を耐支持体上 に含有する、画像的に露光されたハロゲン化模材 を次の処理段階:

(1) 写真材料の現象剤またはその塩の溶液 L による、p H 1~8における1~10秒間の処理、 秒へ短縮することを可能 らしめた (RA4法、コグック2001紙)。

またアダフア95CD型の高級性現像剤の使用 によって具化銀乳剤に基づく常用のカラー紙で同 じ現像時間を達成することも可能になった。

さらに現像時間を無糖することは、黛宝しくぞ して是非考えたいところであるけれども、これは 用いられる現像系における固有の要因によって観 限される。

写真材料が高話性の現像溶液中に浸液されるときこのカラー材料の一番上のカラー層の現像は選ちに開始される。ハロゲン化機の選元の間に放出されるハライドは現像溶液と共に下層にある乳剤層中へ浸透しそしてその現像を妨げる。異なるスペクトル感光性の三つのハロゲン化機乳剤層はこの工程において特に妨害される。

この層の現像の効果は次の三つの要因によって 選組される:

1. 現像剤の拡散過略が一番長い。

(2) 現像剤で含浸された写真材料のアルカリー含有溶液 I による、p H I O ~ 1 4 における 1 ~ 1 0 秒間の処理、

に付することから彼る迅速現象方法に関する。

上記した二つの番を用いる全体の現像時間は、かくして45秒よりも考しく短かい。現像剤を含む第一の格においてp H値は低くそのため現像が起らないかまたは小さい程度の現像しか起らした現像が何ら現像工程で阻害されることを可能ならしめる。第二の路はp H値を増大させるためアルカリを含有する; ヒドロキシルイオンの迅速な拡散に基づき現像はすべての層中で同時に 開始される。

現像刑部液 I の p H 値は貯ましくは 3 . 5 ~ 6 . 5 である; 現像刑部液 I のそれは | 2 ~ 1 4 である。

極度に短かい現像時間に加えて、上記した種類の現像器の使用はまた現象剤権被Iの実用的に制限されない安定性に関して利点を有し、なぜなら

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低いpH値において、自物酸化を促進する空気中の酸素の影響は大きい程度に静設されるからである。さらに別の利点は、この迅速運像決が使用する存破I及びIの機能に殆ど完全に無関係であること、及び事実上監波のない操作方式を提供すること、に見出されるべきである。

二つの溶液を用い写真材料を非常に短い時間で 処理する観点からして、進めったローラーまたは 加圧シリンダーによるいわゆる単一適用を実施す ることもまた可能である。

アルカリ帯として何えばNaOH、KOH、焼 漱三カリウムまたはこれら物質の混合物を用いる ことができる。

成る場合には、ハロゲン化銀の溶解液に影響を 及ぼす物質例えばKCl、KBr、Klまたは安 定剤の少量を溶液Iに添加することによって感光 性結果を調整するのが選当である。

本発明方法を連続操作方式で用いる場合、 静液 の乳剤層への浸透を促進しまたはゼラチン及び水 中に存在するカルシウムイオンを達断するところ

1 - ジホスホン酸もまた鉄線化剤として作用する。 鉄端化剤を二つの現像剤溶液に加えることも有 剤である。

特殊の鉄線化剤は例えば 4、5 - ジセドロキシー1、3 - ペンゼンジスルホン酸、5、6 - ジヒドロキシー1、2、4 - ペンゼントリスルホン酸及び3、4、5 - トリヒドロキシー安息香酸である。

カルシウムを競化するには現像群1モル当り好ましくは約0.2~約1.8モルのカルシウム鏡化剤が用いられる。

鉄鎖化剤は現象剤1モル当り約0.02~約0. 2モルの量で用いられる。

特に適当な一級労香族アミノ現像剤はローフエニレンジアミンであり、特にアルキル基及び労者 族核が関係されているかまたは関係されてい いい、N・ジアルキルーローフエニレンジアミンで ある。そのような化合物の例はN・N・ジエチルーローフエニレンジアミン複酸塩、4・N・N・ジェチルーローフエニレンジアミン

(3) の温調剤及び鎖化剤を溶液I及びIへ添加することもまた有利である。

カルシウムイオンを輸化するため適当な輸化剤 は例えばぞれ合体よく知られているアミノポリカ ルポン酸である。そのようなアミノポリカルポン 酸の臭型的な例はニトロトリ酢酸、エチレンジア ミンテトラ酢酸 (BDTA)、1、3ージアミノ ー2ーヒドロキシープロピルテトラ酢酸、ジエチ レントリアミンベンタ酢酸、N・ーピスー(2 ーヒドロキシペンジル)ーエチレンジアミンーN・ ドージ酢酸、ヒドロキシエチルエチレンジアミ ントリ酢酸、シクロヘキサンジアミノテトラ酢酸 及びアミノーマロン酸である。

その他のカルシウム館化剤はポリホスフエート、ホスホン酸、アミノポリホスホン酸及び加水分解されたポリマレイン酸無水物例えばナトリウムへキサメタホスフエート、1-ヒドロキシエタンー1・1-ジホスホン酸及びエチレンジアミンテトラメチレンーホスホン酸である。1-ヒドロキシエタン-1・

塩酸塩、4-(N-エチル-N-2-メタンスルホニルアミノエチル)-2-メチルーローアエニレンジアミンセスキ硫酸塩-水加物、4-(N-エチル-N-2-ヒドロキシエチル)-2-メチルーローフエニレンジアミン硫酸塩及び4-N。N-ジエチルー2.2\*-メタンスルホニルアミノエチルーローフエニレンジアミン塩酸塩である。

野ましくは現像剤は5~100g/a、特に5~30g/aの量で選用される。

更に白色化剤、白カプラー及び抗酸化物質を抵加することが適当でありうる。適当な抗酸化剤は 例えばヒドロキシルアミン及びジェチルヒドロキ シルアミンならびに重視酸塩であり、これらは好ましくは5g/8までの量で用いられる。

適当な付加的成分は光学的光輝剤、滑剤例えば ポリアルキレングリコール、表面沃性剤、安定剤 例えば複素環メルカプト化合物またはニトロペン ズイミダゾール、及び所要のp H 値を形成するた めの剤である。現像剤溶液はまた5g より少ない ペンジルアルコールを含むこともできる:好まし (4)

くはそれはペンジルアルコールを何ら含まない。

上記した迅速現象法に付される写真記録材料の ハロゲン化級乳剤層は塩化級を少くとも80、行 ましくは少くとも95モル%合有すべきである。

シアンカプラーを含む層は通常赤底光性であり、 マゼンタカプラーを含む層は通常緑底光性であり そして質カプラーを含む層は通常脊膜光性である。

迅速現像法に適当な温度は20~40℃の範囲内にある。

直でに使用できる療液は各個成分から、または 各個成分が相当高機度に溶けているいわゆる 製することができる。機化物は、いできる できる機関が直ぐに使用を使用が直でに使用が変更が直で、それかった。 直では、そのために対象して、の補充ではがませる。 できる。この補充ではがませる。 では、そのためには原にあれて、または他方ではがませるにはできる。 できるにはなったが表別してものでは、 理中に別要されまたは変元の結果としても 現他におけるに関係して特出されるのは 現他に別要された対針に随件して特出される化学

### **実 施 何**

比較の残像:

塩化物乳剤に基づく市駅の常用カラー紙、例えばコダック2001またはアグフアカラー・タイプ9をRA4法の仕様に従い現像し、次いで銀白ー定着しそして洗滌する。

処理は次のようにして行なわれた:

現像:

45 10 . 35 TO

蒙白 - 定着格:

45%, 35%

洗滌:

908.300

老猫.

裕粗成

現像路:

トリエタノールアミノ

11.0 =4

N.N-ジェテルヒドロキシル

アミン

5 - 1 0

4-アミノ-N-エチル-N(β-

メタンスルホンアミドエチル)-

n-トルイジン-セスキ破離塩-

水化物

5.09

使用中の現象剤溶液に補填するために使用することができる。クロライドイオンは通常新たに創製される現像剤の場合を除いては添加することは要せず、それはクロライドイオンは通常現像の結果として写真材料から遊離されるからである。

本発明による現象者の助けにより、その感光性 乳剤層が少くとも80モル%の塩化物を含むハロ ゲン化銀粒子を含有するカラー写真ネガ紙を20 砂以内に最大40℃の風度で現象することが可能 であり、生成される細像はすぐれた品質を有しそ して同じカラーネガ紙からRA-4後により45 砂で得られるものと同等である。特にカラーの光 安定性に関して何ら譲ることがない。

全体の現像時間は好ましくは10秒以下である。 現像の後写真材料は常法により停止され、間白 され、定者され、洗練されそして乾燥されるが、 源白 - 定着操作中で漂白及び定着を続けそして洗 源工程を安定化格で置きかえることが可能である。 また蛋白俗または原白 - 定着浴が十分に酸性であ るならば停止浴を省略することもできる。

塩化カリウム

2.39

エチレンジアミンテトラ酢機

3.00

3.4-ジヒドロキシ-1.2.5-ペンゼン-

トリスルホン酸、

三ナトリウム塩

0.60

・炭酸カリウム

25.09

その他常用の表面活性剤及び

光学的光辉剂。

水を加えて14 とする、pH-10.04

漂白-定着格:

ナトリウムジサルフアイト

159

アンモニウムチオサルフェート 100g

アンモニウム鉄エチレン

ジアミンテトラ酢酸

5 O ø

エチレンジアミンテトラ酢酸

5 ,

混合物も10 とする、 p H - 8.0

現象のみに 4 5 秒を要した処理後、次の最大密度が得られる:

黄 243

マゼンタ 247

シアン 249

## 本発明による現像

. . .

上記比較試験におけると同じ写真材料と用いる ; RA-4現像で処理する代りに、先ず5 間、 30でで次の前波1を用いて処理する:

ナトリウムサルフアイト

4-アミノ-N-エチル-N-(β-

メタンスルホンアミドエチル)-

m-トルイジンセスキ硬酸塩一水化物 20g 水を加えて1,000mgとする、pH=5.0 次に材料を次配組成の溶液 1に3秒間、30℃ で付する:

燐酸三カリウム

509

水酸化カリウムでpHI3に調製

水で1.000mgとする、pH-13

次に停止を2秒間行ないそして常法により処理 を続ける。

最大密度:黄

240

マゼンタ

265

シアン

2 4 7

乳剤層が少くとも 8 0 モル%の塩化物を含有する 上記館 1 項記載の方法。

- 3、現象されるべき写真材料のハロゲン化銀乳剤 層が少なくとも95モル%の塩化物を含有する上 記第1項記載の方法。
- 4. 現像が最大 2 0 秒間内に完了される上記第 1 項記載の方法。
- 5. 現象が最大 1 0 秒 間内に完了される上記第 1 項記載の方法。
- 6. 現象削溶液 I がペンジルアルコーを含まない 上記第1項記載の方法。
- 7. 二つ現象格中において現像が20~40℃範囲内の温度で行われる上記第1項記載の方法。
- 8. 現象剤が5~100g /4 の量で避用される 上記第1項記載の方法。

特許出職人 アグファーゲヴェルト・ アクチェンゲゼルシャフト

代 粟 人 弁理士 小田島 平 吉



(5) 二つの現像工程の比較は、本発明による現象が 値か8秒間の全体現象時間中に所要の最大密度を 有効に生ますること及び得られる値がRA-4独 による値よりも或る程度高いことを明らかに示している。

本発明の主なる特徴及び無様は以下の通りであ ス

- 1.シアンカプラー、マゼンタカプラー及び食力 プラーがそれぞれスペクトル的に適合されている 少くとも三つの異なるスペクトル感光性を有する 感光性ハロゲン化機乳剤腫を層支持体上に含有す る、固像的に露光されたハロゲン化級記録材料を 次の処理股階:
- (1) 写真材料の現像剤またはその塩の痞液 I による、p H l ~ 8 における l ~ 1 0 秒間の処理、
- (2) 現象割で含硬された写真材料のアルカリー含有溶液 R による、 p H 10~14 における 1~10 秒回の処理、

に付することから成る迅速現象方法。

2. 現像されるべき写真配録材料のハロゲン化銀

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(71)Applicant: KONICA CORP

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**UEDA YUTAKA** KOBAYASHI HIROAKI **HAGIWARA MOEKO** 

(54) METHOD FOR PROCESSING SILVER HALIDE PHOTOGRAPHIC SENSITIVE MATERIAL AND DEVELOPER **USED THEREFOR** 

(57)Abstract:

PROBLEM TO BE SOLVED: To provide a silver halide photographic sensitive material processing method by capable of obtaining a high density image having low fog rapidly even at the time of supplying the a small amount of developer with a dry feeling similar to that in a dry process and being stably and rapidly processed without causing any precipitation in a processing solution even at the time of processing a small amount of the sensitive material and even under any circumstances.

SOLUTION: This developer consists of at least two processing solutions, namely a concentrated processing solution having a pH of ≤7 and another concentrated processing solution having a pH of ≥8 and respectively prescribed amounts of these solutions are supplied to the image forming surface of the sensitive material to develop the sensitive material. Also, the developer consists of at least two processing solutions, namely a concentrated processing solution having a pH of ≤7 and another concentrated processing solution having a pH of ≥ 8 and respectively prescribed amounts of these processing solutions are supplied to the image forming surface of the sensitive material to directly mix the processing solutions together on and/or within the photographic image forming layers.

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### **CLAIMS**

#### [Claim(s)]

[Claim 1] The art of the silver-halide photosensitive material characterized by for the processing liquid for development consisting of at least two, pH seven or less strong solution and pH eight or more strong solutions, and carrying out specified quantity supply and developing each processing liquid for development to the image formation side of silver-halide photosensitive material.

[Claim 2] The art of a silver-halide photosensitive material according to claim 1 to which the amount of supply of seven or less above [ pH ] liquid and pH eight or more liquid is characterized by being 5-150ml per two 1m of silver-halide photosensitive material, respectively.

[Claim 3] The art of a silver-halide photosensitive material according to claim 1 or 2 characterized by seven or less above [pH] liquid containing 0.005-1.00 mols /of color development chief remediest.

[Claim 4] The art of a silver-halide photosensitive material according to claim 1, 2, or 3 characterized by eight or more above [ pH ] liquid containing 0.1-3.5 mols /of alkali chemicalsl.

[Claim 5] The art of a silver-halide photosensitive material according to claim 1, 2, 3, or 4 characterized by supplying either [ at least ] seven or less above [ pH ] liquid or pH eight or more liquid to the image formation side of silver-halide photosensitive material through space.

[Claim 6] The art of a silver-halide photosensitive material according to claim 5 characterized by supplying the both sides of seven or less above [pH] liquid and pH eight or more liquid to the image formation side of silver-halide photosensitive material through space.

[Claim 7] The art of a silver-halide photosensitive material according to claim 1, 2, 3, 4, 5, or 6 to which one liquid is characterized by the ratio of the capacity of seven or less above [ which is supplied to the image formation side of silver-halide photosensitive material / pH ] liquid and pH eight or more liquid being less than 100 times to the liquid of another side.

[Claim 8] The art of a silver-halide photosensitive material according to claim 1, 2, 3, 4, 5, 6, or 7 characterized by time after the 1st liquid of the processing liquid for development is supplied to the image formation side of silver-halide photosensitive material until this sensitive material results in the following down stream processing being 5 – 45 seconds.

[Claim 9] The development agent of the silver-halide photosensitive material which consists of at least two, pH seven or less strong solution and pH eight or more strong solutions, and specified quantity supply is carried out in the image formation side of silver-halide photosensitive material, respectively, and is on a photograph composition layer and/or among a photograph composition layer, and is characterized by mixing directly.

[Claim 10] The development agent of a silver-halide photosensitive material according to claim 9 characterized by seven or less above [ pH ] liquid containing 0.005-1.00 mols /of color development chief remediesl. [Claim 11] The development agent of a silver-halide photosensitive material according to claim 9 or 10 characterized by eight or more above [ pH ] liquid containing 0.1-3.5 mols /of alkali chemicalsl.

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#### DETAILED DESCRIPTION

[Detailed Description of the Invention] [0001]

[Industrial Application] About the development agent used for the art of silver-halide photosensitive material, and it, this invention is quick processing of dry feeling in detail, and the picture of high concentration and low fogging is acquired, and it relates to the development agent used for the art of silver-halide photosensitive material and it which are excellent also in the shelf life of processing liquid.

[0002]

[Description of the Prior Art] In the field of processing of silver-halide color photography sensitive material (henceforth sensitive material), the request of speeding up of processing is increasing increasingly.

[0003] On the other hand, since it corresponds to waste fluid regulation in recent years, also in the so-called minilaboratory, low supplement-ization is advancing. For this reason, in a mini-laboratory with few throughputs per day, it is in the inclination for the update rate of processing liquid to fall, especially the color development liquid high—concentration—ized for quick processings tends to receive air oxidation, and it is easy to produce the problem of it becoming impossible to maintain the deposit of sedimentation, generating of tar, and the processability ability further stabilized by degradation of liquid. On the other hand, although mode of processing of the method which encloses color development liquid with the good container of sealing nature, and is sprayed is indicated by JP,6–324455,A in order to prevent air oxidation, development property sufficient with this technology is not acquired, but there is no \*\*\*\*\* in practical use.

#### [0004]

[Problem(s) to be Solved by the Invention] By the way, a mini-laboratory store increases quickly in recent years, and waste fluid is not generated, but an unfamiliar person also has the needs for the processing system which can be used easily in the treatment of the device of the dry feeling in which it does not impress using processing liquid. Moreover, no matter it may install some outdoors, such as change of atmospheric temperature, etc. in what environment, a system which can be processed stably is also desired.

[0005] Although it has succeeded in the attempt which supplies a developer to the image formation side (it is also called an emulsion side) of sensitive material directly, and keeps developer composition constant in order to acquire processing stability The well-known direct supplying method is what supplies comparatively a lot of developers to an emulsion side. Not all the developers to supply permeate into sensitive material, liquid sagging occurs, therefore the amount of supply of processing liquid needs to be little to the grade which does not start liquid sagging, when aiming at dry feeling also at processing stability.

[0006] However, it is difficult to elute halogenide ion from sensitive material, to become high concentration especially if there is little processing liquid, to suppress advance of a development reaction as silver development advances, when it is color development processing, and to obtain required concentration by short—time processing, and it is difficult to use the color development liquid used now as it is.

[0007] Although it is possible to warm sensitive material in order to promote a development reaction, if an elevated temperature is applied from the exterior, fogging will occur. Moreover, even if it warms processing liquid, temperature of sensitive material cannot be enough raised by the few amount of supply. Furthermore, if processing liquid is warmed, oxidization of a developing agent will tend to take place, and generating of tar or precipitation will be caused.

[0008] Even if this invention is made in view of the above-mentioned situation and the purpose makes [ 1st ] a dev loper little supply of dry feeling, it is quick and the processing from which the picture of high concentration and low fogging is acquired is possible, and it is in offering the art of the silver-halide photosensitive material which does not produce the problem of precipitation generating of processing liquid. Moreover, it is in providing the 2nd with the art of the silver-halide photosensitive material which can perform stable quick processing no matter it may be processed in what environment, even if a throughput is little.

[Means for Solving the Problem] As for the abov -mentioned purpose of this invention, the processing liquid for \*\* d velopment consists of at least two, pH seven or less strong solution and pH eight or more strong solutions. The art of the silver-halide photosensitive mat rial which carries out specified quantity supply and develops each processing liquid for development to the image formation side of silver-halide photosensitive material, The amount of supply of seven or less above [ pH ] liquid and pH eight or more liquid is 5-150ml per two 1m of silver-halide photosensitive material, respectively, S ven or less above [ pH ] liquid contains 0.005-1.00 mols /of color

development chief remediesl., Eight or more above [ pH ] liquid contains 0.1–3.5 mols /of alkali chemicalsl., Either [ at least ] seven or less above [ pH ] liquid or pH ight or more liquid is supplied to the image formation side of silver—halide photosensitiv material through space, The both sides of seven or less abov [ pH ] liquid and pH eight or more liquid are supplied to the image formation side of silver—halide photosensitive material through space, The ratio of the capacity of seven or I ss above [ which is supplied to the image formation side of silver—halide photosensitive material / pH ] liquid and pH eight or mor liquid is [ one liquid ] less than 100 times to the liquid of another side, Time after the 1st liquid of the processing liquid for development is supplied to the image formation side of silver—halide photosensitive material until this sensitive material results in the following down stream processing is 5 – 45 seconds, And it consists of at least two, a with a \*\* pH of seven or less strong solution and pH eight or more strong solutions. Specified quantity supply is carried out in the image formation side of silver—halide photosensitive material, respectively. The development agent of the silver—halide photosensitive material which is on a photograph composition layer and/or among a photograph composition layer, and is mixed directly, and seven or less above [ pH ] liquid contain 0.005–1.00 mols /of color development chief remedies!., It is attained more by that eight or more above [ pH ] liquid contains 0.1–3.5 mols /of alkali chemicals!

[0010] That is, this invention person used to divide the processing liquid for development into a strong solution with low pH, and a strong solution with high pH, he used to supply each liquid to the image formation side of sensitive material, regulating an amount, used to think that a development reaction will be promoted using the heat of neutralization which generates both liquid by being on the image formation side of sensitive material, and/or among a photograph composition layer, and mixing directly, and used to result in this invention. When it is an object for the color developments for color sensitive material, as for the processing liquid for development, it is effective to make low pH liquid contain a color development chief remedy by the concentration of 0.005–1.00 mols/l., to high—concentration—ize, to make high pH liquid contain alkali chemicals by the concentration of 0.1–3.5 mols/l., and to be referred to as high concentration and high pH.

[0011] Now, consider as 2 liquid system divided into the solution of low pH which contains a color development chief remedy for color development liquid, and the solution containing alkali chemicals of high pH, and sensitive material is made to flood with each liquid one by one, or liquid is given to JP,2-203338,A with a roller, the permeability of a color development chief remedy is raised, and quickening processing of a color paper is indicated. However, since time is set and alkali chemicals are supplied after supplying a color development chief remedy, this method is in the start of a development reaction, and it is inadequate. [ of the effect of development promotion ] [0012] Hereafter, this invention is explained in full detail for every item.

[0013] [Silver-halide photosensitive material] As an example of the sensitive material processed by the method and processing agent of this invention, the silver-halide color photography sensitive material containing chloride emulsion and the silver-halide color photography sensitive material containing iodine silver bromide or a silver-bromide emulsion are mentioned. Generally such sensitive material comes to prepare the photograph composition layer which consists of hydrophilic-colloid layers, such as photosensitive emulsion layers, such as a blue-sensitive silver-halide emulsion layer, a green sensibility silver-halide emulsion layer, and a red-sensitive silver-halide emulsion layer, an antihalation layer, an interlayer and a filter layer, and a protective layer, on the base material which makes a resin and paper a base. The development agents of this invention are on these photograph composition layers and/or among a photograph composition layer, and it is mixed directly and they function. [0014] [Processing liquid supply] In this invention, it is desirable to supply either [ at least ] pH seven or less strong solution or pH eight or more strong solutions through space. It says making not a form like processing by the usual auto-processor by minding space that makes sensitive material completely immersed in the bath of processing liquid but processing liquid fly here to the image formation side of sensitive material, or applying processing liquid to it using a curtain coating machine or sponge.

[0015] A processing liquid flight means to make processing liquid fly through space as a concrete processing liquid supply means to sensitive material, a processing liquid application means to apply processing liquid to sensitive material through space like a curtain coating machine, etc. are mentioned. As a processing liquid flight means to make processing liquid fly through space to sensitive material Like the thing of the same structure as the ink-jet head section of an ink jet printer, or the thing of structure given in JP,6-324455,A The thing which makes sensitive material generate [ in / the flight means after processing / for processing liquid ] a pressure through space, and makes it fly actively, the thing which makes processing liquid fly like a spray bar according to the fluid-pressure force applied to the flight means after processing through space at sensitive material are mentioned. Since what supplies processing liquid by vibration, the thing which supplies processing liquid by bumping are mentioned as a processing liquid flight means to make the processing liquid by the thing of the same structure as the ink-jet head section of an ink jet printer fly through space to sensitive material, the processing liquid amount of supply is controlled and the processing position of Japanese lacquer and sensitive material can also be chosen, it is desirable.

[0016] The liquid supplied through space may fluctuate an amount according to a supply position. Moreover, it is desirable that both pH seven or less strong solution and pH ight or more strong solutions are preferably supplied through space.

[0017] Moreover, the thing by which the thing which supplies processing liquid to s nsitiv material through space as a processing liquid supply means from a linear supply head, or the thing which supplies processing liquid to sensitive material through space from a field-like supply head also supplies processing liquid to sensitive material through space from a punctiform supply head, or the oth r methods may be used. When sensitive material is a

sheet, in moreover, the state where the physical relationship of sensitive mat rial and a supply head is being fixed, using the supply head of the shape of a field equivalent to the size of sensitive material Although processing liquid may be supplied to sensitive material through space from a supply head, ven if it is [a supply head] smaller to supply processing liquid to sensitive material through space from a supply head, shifting the physical relationship of a supply head and s nsitive material, processing liquid can fully be supplied to sensitive material, and it is desirable. Moreover, in order to supply processing liquid to sensitive material quickly although a supply head may move when using a linear supply head, it is desirable to move sensitive material to a linear supply head in addition to a linear supply head and a linear parallel direction. In order to make the processing time regularity especially, it is desirable to move sensitive material to a linear supply head and a linear perpendicular direction.

[0018] The amount of supply in this invention is processing volume supplied to a direct emulsion side, when it minds space, and when making sensitive material immersed and supplying processing liquid, it points out the amount of supplements of a supplement agent. As for the liquid which contains a color development chief remedy at least, it is desirable that the emulsion side of sensitive material is supplied directly.

[0019] [Development process] The processing liquid for development points out all the liquid containing a compound with the capacity which can contribute to the development reaction of sensitive material, for example, a color development chief remedy, alkali chemicals, etc. by this invention. For example, solution, water, etc. containing a surfactant, the solubilizing agent of a color development chief remedy, preservatives, etc. are included. In this invention, although it consists of at least two processing liquid, pH seven or less strong solution and pH eight or more strong solutions, as for both, it is still more desirable that it is pH 4 or less and pH 10 or more.

[0020] A development process says a thing after supplying the liquid for the first development to sensitive material until it supplies the processing liquid (for example, a bleach fix bath, bleach liquor, the stop solution, etc.) of the following process, or until it floods with the processing liquid of the following process by this invention. Moreover, time to pass a development process is time after supplying the liquid for the first development to sensitive material until it supplies the processing liquid of the following process, or until it floods with the processing liquid of the following process, and is 5-20 seconds preferably about 5 to 45 seconds.

[0021] As for the ratio of the amount of supply of pH seven or less liquid and pH eight or more liquid, it is desirable that one side is less than 100-time capacity to another side, and it is less than double precision further less than 10 times more preferably. About 5-150ml per two of amounts to which each is supplied is 10-100ml and further 10-50ml preferably 1m of sensitive material, and about 10-300ml per two of sum total amount of supply of all processing liquid is 10-100ml and further 20-60ml preferably 1m of sensitive material.

[0022] As time when the processing liquid for development is altogether supplied on the image formation side (henceforth an emulsion side) of sensitive material, it is [less than 2/3 of the beginning that passes a development process] desirable that it is less than 1/3 and less than 1/10 more preferably.

[0023] Although it is desirable to make it be proportional to the light exposure to sensitive material as for supply of the processing liquid for development, it is not necessary to make it be not necessarily proportional. Moreover, in the case of color photography sensitive material, the following desirable examples are given as sequence of supply of each part liquid separation.

[0024] (1) color development chief-remedy content liquid -> — alkali-chemicals content liquid (2) color development chief-remedy content liquid -> alkali chemicals and color development chief-remedy content liquid (3) water -> color development chief-remedy content liquid -> — alkali-chemicals content liquid (4) Water -> color development chief-remedy content liquid -> alkali chemicals and color development chief-remedy content liquid (5) alkali-chemicals content liquid -> — color development chief-remedy content liquid (6) alkali chemicals and color development chief-remedy content liquid (7) Water -> alkali-chemicals content liquid -> color development chief-remedy content liquid (8) In color development chief-remedy content \*\*\*\*, as a desirable example Water -> alkali chemicals and color development chief-remedy content liquid -> [(1), ](2), (3), and (4) are mentioned and it is (1) and (3) still more preferably.

[0025] As for a color development chief remedy, it is desirable that it is the p-phenylene diamine system compound which has a water-soluble machine. What has the aforementioned water-soluble machine on [ at least one ] the amino group of a p-phenylene diamine system compound or a benzene nucleus is mentioned. As a concrete water-soluble machine, it is -(CH2) n-CH2OH, -(CH2) m-NHSO2-(CH2) nCH3, -(CH2) m-O-(CH2) n-CH3, and - (CH2CH2O) nCmH2m+1 (m and n express zero or more integers, respectively.). -COOH basis, -SO3H set, etc. are mentioned as a desirable thing.

[0026] As an example of the concrete compound preferably used by such color development chief remedy, following (C-1) - (C-18) is mentioned.

[0027]

[Formula 1]

# [例示発色現像主薬]

$$(C-1) \qquad \qquad (C-2)$$

$$(C-3) \qquad \qquad (C-4)$$

$$(C-5) \qquad (C-6)$$

$$(C-7) \qquad (C-8)$$

[0028] [Formula 2]

[0030] the inside of these compounds -- a desirable thing -- (C-1), (C-2), (C-3), (C-4), and (C-15) -- and (C-17) (C-18) -- it is .

[0031] In addition, the color development chief remedy used preferably is a p phenylenediamine system color

development chief remedy which has the water-soluble machin expr ssed with the following general formula [P]. [0032]

$$R_3 - N - R_4 - NHSO_2 - R_5$$

$$R_2 \longrightarrow R_1$$

$$NH_2$$

[0033] (In a general formula [P], R1 and R2 express a hydrogen atom, a halogen atom, an alkyl group, an alkoxy group, or the acylamino machine.) R3 expresses an alkyl group and R4 expresses an alkylene machine. R5 expresses the alkyl group or aryl group which is not replaced [ substitution or ].

As an example of these concrete compounds, following compound (C-19) - (C-35) is mentioned. These compounds are shown by the concrete basis of R1-R5 of a general formula [P] being shown. [0034]

[Formula 5]

į, ormana	R <sub>1</sub>	R 2	R <sub>3</sub>	R 4	R <sub>5</sub>
C-19	-н	-н	—C₃H,	-CH2CH(-CH3)-	CH₃
C-20	NHCOCH ₃	-н	—CH₃	-CH2CH2-	-CH <sub>3</sub>
C-21	<b>—н</b>	—H	—CH₃	-CH2CH(-CH3)-	— CH <sub>3</sub>
C-22	—CH₂CH₃	$-\mathbf{H}$	—CH₃	-CH2CH2-	CH₃
C -23	−CH³	—н	—CH₃	-CH₂CH(-CH₃)-	— CH₂CH₃
C-24	−CH³	-н	—CH₃	-CH2CH2-	-CH2CH3
C-25	-0-CH <sub>2</sub> CH <sub>3</sub>	-н	—CH₂CH₃	-CH(-CH <sub>3</sub> )CH <sub>2</sub> -	-CH3
C-26	NHCOCH ₃	—н	-C <sub>3</sub> H <sub>7</sub>	-CH2CH2-	-CH <sub>3</sub>
C-27	−CH³	—н	—CH₂CH₃	- CH2CH2-	-CH <sub>2</sub> -0-CH <sub>3</sub>
C-28	—н	—н	—CH₃	-CH2CH2-	$-CH_2-N-(CH_3)_2$
C-29	−CH₃	—н	—CH₂CH₃	-CH₂CH₂-	-CH <sub>2</sub> CI
C-30	−CH a	—н	—CH₂CH₃	-CH2CH2-	-CH <sub>2</sub> -NHCO-CH <sub>3</sub>
C-31	—CH₂CH₃	—н	—CH₂CH₃	- CH <sub>2</sub> CH <sub>2</sub> -	-CH <sub>2</sub> -0-CH <sub>3</sub>
C -32	—CH;	—н	—CH₂CH₃	- CH <sub>2</sub> CH <sub>2</sub> -	-CH <sub>2</sub> -0-CH <sub>2</sub> CH <sub>3</sub>
C - 33	−CH₃	—н	—CH₂CH₃	— CH₂CH₂CH₂—	-CH <sub>3</sub>
C -34	-cı	—н	—CH₃	- CH2CH2CH2-	-CH3
C -35	-0-CH <sub>3</sub>	—н	-CH2CH3	-CH₂CH(-CH₃)-	-CH <sub>9</sub>

[0035] Among the above-mentioned instantiation compounds, preferably, it is (C-20), (C-27), (C-29), (C-30), and (C-33), and is desirable most in all instantiation compounds (C-1). Moreover, the compound of a general formula [P]

is compoundable according to the method of a publication to JP,4-37198,A. The above-mentioned color development chief remedy is usually used in the form of salts, such as a hydrochloride, a sulfate, and a p-toluenesulfonic-acid salt.

[0036] Moreover, the aforementioned color development chief remedy is independent, or it may be used together two or more sorts, and may use together with monochrome developing agent, for example, a phenidone, 4-hydroxymethyl-4-methyl-1-phenyl-3-pyrazolidone, a Metol, etc. by request, and may use.

[0037] About 0.005-1.00 mols /of concentration of the chief remedy of the liquid containing a p phenylenediamine system color development chief remedy are [l.] 0.01-0.25 mols [l.] /and further 0.06-0.13 mols/l. preferably. When this concentration is 0.05 mols/l. or more, as for pH of liquid, four or less are desirable from the soluble point of a chief remedy, and it is pH two or less further. Moreover, when this concentration is 0.25 mols/l. or more, it is d sirable that pH is two or less.

[0038] It is stable in copy genuineness ability to contain the compound shown by the following general formula [H] or [B] in the processing liquid for the color developments, and fogging produced in the unexposed section also has the advantage of being few.

[0039]

[Formula 6] 一般式〔H〕

[0040] In a general formula [H], although an alkyl group, an aryl group, an R'-CO-machine, or a hydrogen atom is expressed, respectively, even if the alkyl group R6 and whose R7 are not hydrogen atoms simultaneously and which is expressed with R6 and R7 is the same, they may differ, and its alkyl group of carbon numbers 1-3 is desirable respectively. Furthermore, these alkyl groups may have a carboxylic-acid machine, a phosphoric-acid machine, a sulfonic group, or a hydroxyl group. R' expresses an alkoxy group, an alkyl group, or an aryl group. Also including that in which the alkyl group and aryl group of R6, R7, and R' have a substituent, it may join together, and R6 and R7 may constitute a ring, for example, they may constitute the heterocycle like a piperidine, a pyridine, triazine, or a morpholine.

[0041] [Formula 7] 一般式 (B)

$$R_8 > N - N < R_{10} < R_{11} = R_{11}$$

[0042] In a general formula [B], R8, R9, and R10 express the alkyl group which is not replaced [ a hydrogen atom, substitution, or ], an aryl group, or a heterocycle machine, and R11 expresses the alkyl group which is not replaced [ a hydroxy group, the hydroxy amino group, substitution, or ], an aryl group, a heterocycle machine, an alkoxy group, an aryloxy group, a carbamoyl group, and the amino group. As a heterocycle machine, it may be 5 - 6 member ring, and saturation or an unsaturation is [ it may consist of C, H, O, N, S, and a halogen atom, and ] sufficient. R12 expresses the divalent basis chosen from -CO-, -SO2-, or -C(=NH)-, and n is 0 or 1. Especially R11 may express an alkyl group, an aryl group, and the basis chosen from a heterocycle machine at the time of n= 0, and R10 and R11 may form a heterocycle machine jointly.

[0043] Especially the compound shown by the following general formula [D] among the compounds of a general formula [H] is desirable.

[0044]

[Formula 8] 一般式〔D〕

$$EO-N$$

[0045] (In a general formula [D], L expresses an alkylene machine, A expresses a carboxyl group, a sulfonic group, a phosphono machine, a phosphinic acid machine, a hydroxyl, the amino group, an ammonio machine, a carbamoyl group, or a sulfamoyl group, and R expresses a hydrogen atom or an alkyl group.) L, A, and R may be replaced by each also in no replacing also including a straight chain and branched chain. L and R may connect and a ring may be formed

It explains still in detail about the compound shown by the general formula [D]. Among a formula, L expresses the alkylene machine which may replace the straight chain of carbon numbers 1–10, or branched chain, and its carbon numbers 1–5 are desirable. Specifically, bases, such as a methylene, ethylene, trimethylene, and a propylene, are mentioned as a desirable example. As a substituent, a carboxyl group, a sulfonic group, a phosphono machine, a

phosphinic acid machine, a hydroxyl, and the ammonio machine that may carry out alkylation are expressed, and a carboxyl group, a sulfonic group, a phosphono machin, and a hydroxyl are mentioned as a desirable xample. A expr sses a carboxyl group, a sulfonic group, a phosphono machine, a phosphinic acid machine, a hydroxyl or the amino group that may carry out alkylation, an ammonio machine, a carbamoyl group, or a sulfamoyl group, and is mentioned as an example with desirable carboxyl group, sulfonic group, hydroxyl, phosphono machine, and carbamoyl group that may carry out alkylation. - As an example of L-A, a carboxymethyl machin, a carboxy ethyl group, a carboxy propyl group, a sulfoethyl machine, a sulfo propyl group, a sulfo butyl, a phosphono methyl group, a phosphono ethyl group, and a hydroxyethyl machine can be mentioned as a desirable example, and a carboxymethyl machine, a carboxy ethyl group, a sulfoethyl machine, a sulfo propyl group, a phosphono methyl group, and a phosphono ethyl group can be especially mentioned as a desirable example. R expresses the alkyl group which may replace the straight chain of a hydrogen atom and carbon numbers 1-10, or branched chain, and its carbon numbers 1-5 are desirable. As a substituent, a carboxyl group, a sulfonic group, a phosphono machine, a phosphinic acid machine, a hydroxyl or the amino group that may carry out alkylation, an ammonio machine, a carbamoyl group, or a sulfamoyl group is expressed. There may be two or more substituents. As R, a hydrogen atom, a carboxymethyl machine, a carboxy ethyl group, a carboxy propyl group, a sulfoethyl machine, a sulfo propyl group, a sulfo butyl, a phosphono methyl group, a phosphono ethyl group, and a hydroxyethyl machine can mention as a desirable example, and can mention as an example with especially desirable hydrogen atom, carboxymethyl machine, carboxy ethyl group, sulfoethyl machine, sulfo propyl group, phosphono methyl group, and phosphono ethyl group. L and R may connect and a ring may be formed.

[0046] Next, although the typical example of a compound is shown among the compounds expressed with a general formula [D], this invention is not limited to these compounds.

[0047]

[Formula 9]

[0048] [Formula 10]

[0049] [Formula 11]

[Formula 12]

[0051] The compound shown by these general formulas [H] or the general formula [B] is usually used in forms, such as the amine of isolation, a hydrochloride, a sulfate, a p-toluenesulfonic-acid salt, an oxalate, phosphate, and acetate.

[0052] A sulfite can be used for the processing liquid for the color developments as preservatives. As this sulfite, a sodium sulfite, potassium bisulfite, etc. are mentioned. That may not be right although \*\*'s of a sulfite contained in the same partial liquid as a color development chief remedy is desirable. The concentration of a sulfite has desirable l. in  $1\times10^{-4}$  to  $5\times10^{-4}$  to

[0053] A buffer can be used for the processing liquid for the color developments, as a buffer Potassium carbonate, a sodium carbonate, a sodium bicarbonate, a potassium bicarbonate, Phosphoric-acid 3 sodium, phosphoric-acid 2 potassium, the sodium borate, A boric-acid potassium, tetraboric-acid sodium (boric acid), a tetraboric-acid potassium, Ortho-hydroxyb nzoic-acid sodium (sodium salicylate), an ortho-

hydroxybenzoic-acid potassium, A 5-sulfo-2-hydroxy sodium benzoate (5-sodium sulfosalicylate) and a 5-sulfo-2-hydroxybenzoic-acid potassium (5-sulfosalicylic-acid potassium) are desirable.

[0054] Alkali chemicals are used for the processing liquid for the color developments, and a lithium hydroxide, a sodium hydroxide, a potassium hydroxide, etc. ar mentioned other than the aforementioned buffer as alkali

chemicals. About 0.1-3.5 mols /of concentration of alkali chemicals are [l.] 0.3-1.2 mols/l. preferably. When alkali chemicals seldom dissolve under the influenc of temperatur or other solutes, it is desirable to use it in the range of the amount which can be dissolved.

[0055] An accelerator can be used for the processing liquid for the color developments, and a thioether syst m compound, a p-phenylene diamine syst m compound, quarternary ammonium salt, para aminophenol, an amine system compound, polyalkylene oxide, 1-phenyl-3-pyrazolidone, hydronalium gin, the Mesoyi-on type compound, an ion type compound, imidazole derivatives, etc. can be added as an accelerator in it if needed.

[0056] As for the processing liquid for the color developments, it is desirable not to contain benzyl alcohol substantially.

[0057] In the processing liquid for the color developments, a chloride ion and bromine ion can be added for the purpose, such as fogging prevention. When directly added by color development liquid, although the chloride of sodium, a potassium, ammonium, nickel, magnesium, manganese, calcium, or cadmium is mentioned, desirable things are a sodium chloride and potassium chloride as chloride-ion feed materials. Moreover, it may be supplied with the form of the opposite salt of the fluorescent brightener added by color development liquid. As feed materials of bromine ion, although the bromide of sodium, a potassium, ammonium, a lithium, calcium, magnesium, manganese, nickel, cadmium, a cerium, or a thallium is mentioned, desirable things are a potassium bromide and a sodium bromide. Not containing substantially is most desirable although at most 0.02 mols /of contents of these halogen ion are [ I. ] 0.001 mols/l. or less preferably.

[0058] It is desirable to make the processing liquid for the color developments contain a thoriadinyl stilbene system fluorescent brightener, and the compound specifically shown by the following general formula [E] is desirable. [0059]

[Formula 13] 一般式〔E〕

[0060] Setting at an upper ceremony, X2, X3, Y1, and Y2 are halogen atoms, such as a hydroxyl group, chlorine, or a bromine, an alkyl group, an aryl group, and [0061] respectively.

[Formula 14]

$$-N \stackrel{R_{15}}{\underset{R_{14}}{\stackrel{R_{15}}{\longrightarrow}}} 0$$

[0062] Or -OR17 is expressed. Respectively, it is here, and in R13 and R14, R15 and R16 express an alkylene machine (a substitution product is included), R17 expresses a hydrogen atom, an alkyl group (a substitution product is included), or an aryl group (a substitution product is included) for a hydrogen atom, an alkyl group (a substitution product is included), and M expresses a cation.

[0063] Furthermore, in addition to this, various additives, such as a stain inhibitor, a sludge inhibitor, and an

interlayer effect accelerator, can be used again.
[0064] Moreover, it is desirable that the chelating agent shown in the processing liquid for the color developments

[UU64] Moreover, it is desirable that the chelating agent shown in the processing liquid for the color developments by following general formula [K-I]-[K-V] is added. [0065]

$$\begin{array}{c} \text{M}_{1} \text{OOC} - \text{CH}_{2} \\ \text{M}_{2} \text{OOC} - \text{CH}_{2} \end{array} > N - \begin{array}{c} \text{A}_{3} & \text{A}_{1} \\ \text{I} & \text{I} \\ \text{C} - \text{CH} \\ \text{I} & \text{I} \\ \text{A}_{4} & \text{A}_{2} \end{array}$$

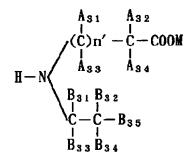
$$A_{11}$$
 - CHNH - X - NHCH -  $A_{13}$   
 $A_{12}$  - CH<sub>2</sub> CH<sub>2</sub> -  $A_{14}$ 

$$\begin{array}{c} A_{21} - (CH_2)n_1 \\ A_{22} - (CH_2)n_2 \end{array} N - X_1 - N \left( \begin{array}{c} (CH_2) \\ \hline \\ (CH_2) \\ \hline \\ \hline \\ \end{array} A_{24} \end{array} \right)$$

[0068]

[Formula 18]

一般式 [K-N]



[0069]

[Formula 19]

一般式 [K-V]

[0070]

[Formula 20]

$$K - I - 1$$
 OH
 $HOOC - CH_2 > N - CH_2 - CH - COOH$ 

$$K - I - 2$$
 $H00C - CH_2 > N - CH_2CH_2C00H$ 

$$K - I - 5$$
 $HOOC - CH_2 > N - CH_2 CHCOOH$ 
 $CH_2 COOH$ 

[0071] [Formula 21]

 $\begin{array}{lll} {\rm H}_2{\rm O}_3{\rm P-CH-NH-CH}_2{\rm CH}_2{\rm CH}_2{\rm -NH-CH-PO}_3{\rm H}_2 \\ {\rm H}_2{\rm O}_3{\rm P-CH}_2 & {\rm CH}_2{\rm PO}_3{\rm H}_2 \end{array}$ 

http://www4.ipdl.jpo.go.jp/cgi-bin/tran\_web\_cgi\_ejje

K - II - 9

[0072] [Formula 22]

[0073] [Formula 23]

$$K - m - 10$$

$$\begin{array}{c|c} \operatorname{HOOC-CH_2} & \operatorname{N-CH_2CH_2NCH_2CH_2-N} & \begin{array}{c} \operatorname{CH_2-COOH} \\ \operatorname{CH_2-COOH} \end{array} \\ \end{array}$$

$$K - N - 1$$

$$K - N - 2$$

$$K-V-1$$
 0  $CH_3$  0  $H0-P-C-P-OH$  0  $H$  0  $H$  0  $H$ 

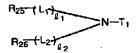
$$K-V-2$$
 0  $C_2H_5$  0  $K-V-3$  0 OH OH 0 HO  $-P-C-C-P-OH$  HO  $-P-C-C-P-OH$  OH OH OH OH OH OH

[0074] Also in these chelating agents, it divides and K-I -2, K-II -1, K-II -5, K-III -10, K-IV -1, and K-V -1 are used preferably.

[0075] Furthermore, the processing liquid for the color developments can be made to contain an anion, a cation, both sexes, and the field side activator of a Nonion again, and various surfactants, such as an alkyl sulfonic acid, an aryl sulfonic acid, an aliphatic carboxylic acid, and an aromatic carboxylic acid, may be added if needed. [0076] [Bleaching processing] It is desirable to contain at least one sort of the 2nd iron complex of an amino polycarboxylic acid and hydrate salts in bleaching processing liquid. You may mix and use two or more sorts of different 2nd iron complex of an amino polycarboxylic acid and hydrate salts.

[0077] It is desirable to be used as the 2nd iron complex of an amino polycarboxylic acid as a form of the iron complex of the free acid (compound shown by the following general formula [1]) of an amino polycarboxylic acid shown below, and it is still more desirable to use tog ther the aforem ntioned 2nd iron complex and the free acid of an amino polycarboxylic acid. Especially a desirable thing is using together the aforementioned 2nd iron complex, and constituting it and the free acid of an amino polycarboxylic acid of the same kind. Moreover, the 2nd \*\*\*\*\*\*\*\* hydrate salt of an amino polycarboxylic acid can be used as potassium salt, sodium salt, an ammonium salt, etc., and the fr e acid of an amino polycarboxylic acid can be used as the acid of isolation, potassium salt, sodium salt, etc.

[0078] [Formula 24] 一般式 〔Ⅰ〕



式中、T1は、水素原子、ヒドロキシ基、カルボキシ基、スルホ基、カルバモイル基、ホスホノ基、ホスホン基、スルファモイル基、置換されていても無置換であってもよいアルキル基、アルコキシ基、アルキルスルホンアミド基、アルキルチオ基、アシルアミノ基、ヒドロキサム酸基ヒドロキシアルキル基、又は、

W1は、置換されていても無置換であってもよいアルキレン基、 アリーレン基、アルケニレン基、シクロアルキレン基、アラル キレン基、又は、

[0079] R25-R29 express a hydrogen atom, a hydroxy group, a carboxy group, a sulfonic group, a carbamoyl group, a phosphono machine, a HOSUHON machine, a sulfamoyl group, a sulfone acid machine, the acylamino machine, and a hydronalium KISAMU machine, and at least one of R25-R29 is a carboxy group.

[0080] L1-L7 express the alkylene machine which could be replaced even if replaced, an arylene machine, an alkenylene group, a cyclo alkylene machine, or an aralkylene group. I1-I7 express the integer of 0-6. However, I5-I6 are not in 0 with a bird clapper simultaneously.

[0081] The concrete instantiation compound of an amino polycarboxylic acid (instantiation compound I) shown by the general formula [I] which constitutes the 2nd iron complex of an amino polycarboxylic acid and a hydrate salt is shown below.

[0082]

[Formula 25]

[0083] [Formula 26]

[Formula 27]

[0085] [Formula 28]

[0086] As a desirable compound, – (I-1) (I-8), (I-12), (I-14) – (I-20), (I-22), (I-23), and (I-27) are mentioned, and (I-1), (I-2), (I-3), (I-6), (I-12), (I-14), (I-15), and (I-17) are especially mentioned as a desirable compound.

[0087] The concrete instantiation compound of the 2nd iron complex of an amino polycarboxylic acid and a hydrate salt (instantiation compound II(s)) is shown below.

[0088]

[Formula 29]

アミノホ゛りカルホ゛ン型	アミノポリカルボン酸 Fe(Ⅲ)錯体 例示化合物Ⅱ類							
記号	アミノホ"リカルホ"ン酸 (例示化合物 [ 類)	対カチオン	Feltuc対する 結晶水のモル数					
ו — ו	1-1	Na+	3					
II — 2	,	K+	2					
II — 3	,	NH4+	2					
II — 4	1 — 2	Na+	3					
II — 5	"	K+	1					
П — е	"	NH4+	1					
II —7	1 — 3	K+, H+	1					
II —8	"	NH4+, H+	1					
II —9	T — 5	K+	1					
II —10	"	NH4+	1					
II —11	I —14		2					
II —12	ī —28	K+	1					
II —13	I.—26	. K+	1					
II —14	I —10	_	1.5					
II ─15	I — 8	NH4+	2					

[0089] Moreover, as for bleaching processing liquid, it is desirable to contain the organic-acid compound shown by the following general formula [A].

[0090] General formula [A]

A' expresses the organic machine of n \*\* among an A'(-COOM) n formula, n expresses the integer of 1-6, and M expresses ammonium, alkali metal (sodium, a potassium, lithium, etc.), or a hydrogen atom.

[0091] In a general formula [A], as an organic machine of n \*\* expressed with A' An alkylene machine (a methylene group, an ethylene, a trimethylene machine, tetramethylen machine, etc.), Alkenylene groups (ETENIREN machine etc.), alkynylene group (ethynylene group etc.), Cyclo alkylene machines (1, 4-cyclohexane diyl machine, etc.), arylene machines (o-phenylene group, p-phenylene group, etc.), alkane Trier machines (1, 2, 3-propane Trier machine, etc.), and arenetriyl machines (1, 2, 4-benzene Trier machine, etc.) are mentioned.

[0092] The basis of n \*\* expressed with A' described above contains what has substituents (a hydroxy group, an alkyl group, halogen atom, etc.) (2-hydroxy - 1, 2-dihydroxyethylene, hydroxy ethylene, 1, 2, 3-propane Trier, a methyl-p-phenylene, a 1-hydroxy-2-chloroethylene, a chloro methylene, chloroethenylene, etc.). The desirable example of the compound shown by the general formula [A] below is shown.

[0093]

[Formula 30]

- (A-1) HOOCCH<sub>2</sub>C(OH)(COOH)CH<sub>2</sub>COOH
- (A---2) HOOC(CHOH)<sub>2</sub>COOH
- (A-3) HOOCCH<sub>2</sub>COOH
- (A-4) HOOCCH(OH)CHgCOOH
- $\begin{array}{ccc}
  \text{(A-5)} & \text{H} \\
  \text{HOOC} & \text{C} = \text{C} \\
  \text{H}
  \end{array}$
- (A-6) HOOC C=C COOH
- (A-7) (COOH)2
- (A-8) COOH
- (A-9) COOH
- (A-10)  $0 < CH_2-COOH$
- (A—11) COOH
- (A—12) COOH

[0094] [Formula 31]

(A-13) 
$$HOOC - (CH_2)_3 - COOH$$
  
(A-14)  $HOOC - (CH_2)_4 - COOH$   
(A-15)  $HOOCC = CCOOH$ 

HO --- CH2-

COOH

(A-20)

[0095] Especially a desirable thing in the above instantiation compound An instantiation compound (A-1), It is (A-3), (A-4), (A-5), (A-6), (A-13), (A-14), (A-15), and (A-20), and (A-1), (A-5), (A-6), (A-13), (A-14), and (A-20) are especially desirable. Moreover, as a salt of the aforementioned acid, although an ammonium salt, lithium salt, sodium salt, potassium salt, etc. are mentioned, sodium salt and potassium salt are desirable from a viewpoint of preservation stability. It can also use independently and these organic acid or its salt can also use two or more sorts together.

[0096] Moreover, in bleaching processing liquid, you may contain a re~halogenating agent. Although a thing well–known as a re-halogenating agent can be used, compounds, such as an ammonium bromide, a potassium bromide, a sodium bromide, a sodium bromide, a sodium chloride, an ammonium chloride, a potassium iodide, a sodium iodide, and an ammonium iodide, are mentioned, for example.

[0097]

[Example] Although an example is given and this invention is explained in detail hereafter, the mode of this invention is not limited to this.

[0098] Example 1 <u>drawing 1</u> is the outline block diagram of the principal part of the auto-processor (henceforth an automatic processor) used here. The perspective diagram of the principal part of this auto-processor is shown in <u>drawing 2</u>. Moreover, <u>drawing 3</u> is a perspective diagram near the feed-hopper dryness prevention means of this auto-processor.

[0099] [Equipment] There are the heating drum 11 besides the conveyance roller which is not illustrated, the sticking-by-pressure belt 15, the heating belt 33, a conveyance roller after a bleaching fixing processing tub, etc. as a conveyance means to convey the silver-halide photosensitive material P by the predetermined bearer rate. And it has a sensitive-material detection means 70 to detect existence of sensitive material P in the predetermined position of the conveyance direction upstream of a conveyance means, rather than the point to which the processing liquid supply means 52 supplies processing liquid. And there is a heating means 10 to heat the silverhalide photosensitive material P in the conveyance path lower stream of a river of the sensitive material P of the sensitive-material detection means 70. There is a heating drum 11 in the heating means 10. Moreover, the outlet side roller 12 is in the heating drum 11 bottom. The entrance-side roller 13 is in the left-hand side of the heating drum 11. The sticking-by-pressure belt driving roller 14 is in the entrance-side roller 13 bottom on the left-hand side of the outlet side roller 12. The heating drum 11 is made to stick by pressure and convey sensitive material P by moving, while the outlet side roller 12, the entrance-side roller 13, and the sticking-by-pressure belt driving roller 14 are built over the sticking-by-pressure belt 15, it is brought into the 90-degree section of the peripheral surface of the heating drum 11 and is stuck to the heating drum 11 by pressure. Sensitive material P is heated by these. [0100] The development means 50 is in the conveyance path lower stream of a riv r of the sensitive material P of the heating drum 11. The development means 50 has the first processing liquid container 51 and the second processing liquid container 56 as a proc ssing liquid container which holds the processing liquid (processing liquid for the color developments) which processes sensitive material P. The first processing liquid container 51 and the second processing liquid container 56 ar sealed to the open air. The supply head later mentioned as a processing liquid supply means 52 in this example is used. Thereby, the processing liquid supply means 52 supplies proc ssing

liquid (processing liquid for the color developments) to the emulsion side of the sensitive mat rial P heat d by the heating means 10 through space. Moreover, there is a circulating pump 54 in the left of the second processing liquid container 56 in the upper part of the first processing liquid container 51, a filter 55 is in the second processing liquid container 51, and the processing liquid for the color developments is circulated in the direction shown in the arrow of drawing 1 from the first processing liquid container 51 by operating a circulating pump 54 in order of a circulating pump 54, the second processing liquid container 56, and a filter 55. Moreover, a rotator 57 rotates in the second processing liquid container 56, and stirs the processing liquid in the second processing liquid container 56.

[0101] By this, it will be prepared between the processing liquid supply meanses 52 from the second processing liquid container 56, and will have a filtration means (filter 55) to filter the processing liquid from the second processing liquid container 56. Moreover, a replenisher is supplied to the second processing liquid container 56 from the replenisher supply means 59.

[0102] The first shutter 62 and the second shutter 64 which are stopped in the middle of are formed in the processing liquid supply means 52 in supply of the processing liquid to a supply head. [ the cross direction of sensitive material P ] The first shutter 62 is driven by the first shutter mechanical component 61 free [ insertion secession on the supply way of the processing liquid to a supply head ], and drives the second shutter 64 free [ insertion secession ] on the supply way of the processing liquid to a supply head by the second shutter mechanical component 63. <u>Drawing 2</u> shows the state where the second shutter 64 was inserted in the supply way of the processing liquid to a supply head.

[0103] Under the processing liquid supply means 52, in order to prevent dryness of the processing liquid in the feed hopper of the supply head of the processing liquid supply means 52, when not supplying processing liquid to sensitive material P, the feed-hopper dryness prevention means 80 which covers with the lid of the feed hopper of a supply head is. The feed-hopper dryness prevention means 80 has the movable lid 81, the bearing bar 82 supporting the movable lid 81, and the motor 83 to which a bearing bar 82 is moved up and down. A rack drives a bearing bar 82 up and down by the motor 83 on a motor 83 at a bearing bar 82 preparing a pinion. Although the processing liquid supply means 52 supplies processing liquid periodically also in the standby operating condition which does not process sensitive material P so that the movable lid 81 may be a cross-section concave-like and it may mention later In this case, it prevents that surrounding equipment becomes dirty with this processing liquid by the movable lid's 81 moving downward a little, receiving the processing liquid supplied from the processing liquid supply means 52, letting the hole which was prepared into the bearing bar 82 and which is not illustrated pass, and discharging processing liquid to a waste liquid part.

[0104] A second heating means 30 to heat sensitive material P is in the conveyance path downstream of the sensitive material P of the place to which processing liquid is supplied by the processing liquid supply means 52 through space. There are the heating roller 31, the drive roller 32, and a heating belt 33 in the second heating means 30. The heating roller 31 and the drive roller 32 are built over the heating belt 33. The heating roller 31 is in the conveyance path downstream of the previous sensitive material P with which processing liquid is supplied by the processing liquid supply means 52 through space, and heats the heating belt 33. The drive roller 32 which is in the conveyance path downstream of sensitive material P from the heating roller 31 makes the heating belt 33 drive. This heats sensitive material P, where the heating belt 33 is heated. And the second heating means 30 will heat the silver-halide photosensitive material by which processing liquid was supplied to the emulsion side by the processing liquid supply means 52 through space.

[0105] Then, bleaching fixing processing is carried out with the bleaching fixing processing cistern BF, and stabilizing treatment of the sensitive material P in which color development processing was carried out by the development means 50 is carried out by the stabilizing treatment tub ST.

[0106] The outline block diagram of the principal part of an automatic processor which has the two aforementioned development meanses 50 is shown in <u>drawing 4</u>. In this example, the case where it processes using the automatic processor shown by <u>drawing 1</u> is made into an art 1, and let the case where it processes using the automatic processor shown by <u>drawing 4</u> be an art 2.

[0107] <u>Drawing 5</u> is the schematic diagram of the processing tank part of the automatic processor for immersing development which can supply two kinds of processing liquid for the color developments. After sensitive material P is exposed, they is conveyed with a pair of delivery roller, and is first processed by the development tub CD 1. [ two or more ] It is conveyed one by one by the late-coming color development tub CD 2, the bleaching fixing processing tub BF, and the stabilization tub STB by the roller conveyance means, and is processed, respectively. It dries by the dryer part and the sensitive material P to which these processings of each were performed is discharged outside the plane. Let processing using this automatic processor be an art 3.

[0108] <u>Drawing 6</u> is the schematic diagram of the automatic processor of the type which sensitive material P is made to flood with the processing liquid of the processing tub CD 2, and supplies other partial liquid, after supplying the partial liquid of the processing liquid for the color developments to sensitive material P through space by the processing liquid supply means CD 1. After color development processing, after bleaching fixing processing and stabilizing treatment are performed to sensitive material P, it dries and it is discharged outside the plane. Let processing using this automatic processor be an art 4.

[0109] <u>Drawing 7</u> is the schematic diagram of the automatic processor which can be supplied to the emulsion side of sensitive material P through space, after mixing two kinds of processing liquid for the color developments. Let processing using this automatic processor be an art 5.

[0110] [Processing liquid supply means] A linear supply h ad is used for drawing 1 and the processing liquid supply means of the auto-processor of 4, 6, and 7. This linear supply head is perpendicular to the conveyance direction of sensitive material P. The array of a feed hopper is a staggered arrangement of a biseriate. The interval of a feed hopper is 100 micrometers in the distance between edges with the maximum proximity feed hopper. For the diameter of 100 micrometers (7.85x10 to 9 m area 2) of a feed hopper, the processing liquid amount of supply per two is 25ml in 50ml and arts 2 and 4 at arts 1 and 5 the 5000 number of times of processing liquid supply for 1 second, and 1m of silver-halide photosensitive material.

[0111] [Sensitive material] Konica QA-A, Inc. 6 paper exposed by the usual method is processed.

[0112] [processing liquid prescription: It is ] perl.

<<color development liquid - 1>>

Sodium-sulfite 15.0g4-amino-3-methyl-N-ethyl 0.2g screw (sulfoethyl) hydroxylamine disodium 12.0g diethylenetriamine pentaacetic acid 5 sodium 3.0g polyethylene-glycol #4000 8.0g potassium carbonate 45.0gp-toluenesulfonic acid sodium - N - (beta- (methanesulfon amide))

Ethyl aniline sulfate (CD-3) pH is adjusted to 10.0 using 10.0g potassium hydroxide or a sulfuric acid.

[0113] <<color development liquid - 2>>

Partial liquid A sodium sulfite 0.4g diethylenetriamine pentaacetic acid 5 sodium 3.0g polyethylene-glycol #4000 6.0gp-toluenesulfonic acid sodium 30.0gCD-3 It adjusts to pH given in a table using 40.0g potassium hydroxide or a sulfuric acid.

[0114] Partial liquid B diethylenetriamine pentaacetic acid 5 sodium 3.0g polyethylene-glycol #4000 10.0g potassium carbonate It adjusts to pH given in a table using 90.0g potassium hydroxide or a sulfuric acid.

[0115] <<br/>bleaching fixing and stabilizing treatment process>>

Konica [ Corp. ] make: It carried out using the processing agent for the said processes on the processing conditions of CPK-2-J1 process.

[0116] The color paper was processed for three consecutive weeks at a rate of 2 10m per day using drawing 1 and the auto-processor of 4, 5, 6, and 7. To the art 1, it processed in color development processing-time 10 seconds using color development liquid -1. The replenisher used color development liquid -1 as it was. To arts 2 and 5, it processed in color development processing-time 10 seconds using color development liquid -2. The replenisher used partial liquid A and B as they were. Moreover, at the art 2, the supply interval was supplied in 0.3 seconds in order of partial liquid A and partial liquid B.

[0117]

Processing condition [ of the color development process of the <<art 3 ]>>

Processing liquid supply means Used solution Processing time Processing temperature Amount of supply (second) (degree C) (ml/m2)

CD1 Color development liquid-2 partial liquid A 5 39.5 25 (pH 1.5)

CD2 Color development liquid-2 partial liquid B 5 39.5 25 (pH 13.0)

KCl3.7g/l. addition, in addition the processing time here are time after flooding sensitive material with processing liquid until it floods with the following processing liquid.

[0118]

Processing condition [ of the color development process of the <<art 4 ]>>

Processing liquid supply means Used solution Processing time Processing temperature Amount of supply (second) (degree C) (ml/m2)

CD1 Color development liquid-2 partial liquid A - - 25 (pH 1.5)

CD2 Color development liquid-2 partial liquid B 5 39.5 25 (pH 13.0)

5 seconds after supplying color development liquid-2 partial liquid A from the processing liquid supply means CD 1 through KCl3.7 g/l addition space, you made it immersed in the processing tub CD 2.

[0119] Replenisher>> used with CD1 and CD2 of the <<art 3, and CD2 of an art 4

replenisher for partial liquid A – a sodium sulfite perl. 0.6g diethylenetriamine pentaacetic acid 5 sodium 4.3g polyethylene-glycol #4000 8.6g para toluenesulfonic acid sodium 42.9gCD-3 replenisher for 64.0g partial liquid B – perl. — diethylenetriamine pentaacetic acid 5 sodium 4.3g polyethylene-glycol #4000 14.3g potassium carbonate At the time of 129.0g consecutive-processing start, the sample was processed after one week, two weeks, and three weeks, and the maximum blue reflection density Dmax (Y) and (440nm) were measured. [ Dmin (C), (660nm) and ] [ the part light reflex concentration Dmin of 440nm and 660nm of the unexposed section (Y), (440nm), ]

[0120] Moreover, it saved at the room temperature for two weeks, putting the partial liquid A and B of color development liquid -2 into a processing container or a processing tub, this container and tub after preservation were observed, and the following criteria estimated.

[0121] O: — although \*\*:sludge as which a sludge is not regarded although muddiness of O:liquid with which a sludge is not seen at all is checked is slightly checked by the oil level, x:sludge which is a satisfactory grade is seen considerably

[0122] Furthermore, in the art 2, pH of the partial liquid A and B of color development liquid -2 was changed as shown in Table 1, and it evaluated similarly.

[0123] The above result is shown in Table 1.

[0124]

[Table 1]

実験	処理	完色月 - 2 0			開始時			1 選節後		2 週間後		3 週間後		処理容器。 処理権の様子		主薬 残存率	備考		
No.	方法	部分 液A	部分 液B	Dwax (Y)	Dain (Y)	Dain (C)	Onax (Y)	Dain (Y)	Dmin (C)	Dnax (Y)	Dain (Y)	Omin (C)	Dutax (Y)	Dnin (Y)	Duin (C)	CDI	CD2	(%)	
1	1	+	_	1.23	0.06	0.09	1.21	0.06	0. 10	1.20	0.07	0.10	1. 18	0.08	0.12	_		85	比較例
2	2	1.5	13.0	2.27	0.05	0.06	2.26	0.05	0.06	2.25	0. 05	0.08	2.25	0.05	0.06	0	<b>©</b>	98	本発明
3	3	1.5	13. Q	2, 25	0.05	0.06	2. 98	0.08	0.08	2.06	0. 10	0.10	2.04	0. 12	0. 12	Δ	Δ	92	
4	4	1.5	13.0	2. 25	0.05	0.06	2. 23	0.06	0.07	2.20	0.08	0.09	2. 16	0.07	0.10	<b>©</b>	0	98	
5	5	1.5	13.0	1,98	0.05	0.07	-	–	-		-	-	-	-	-	<b>©</b>	0	98	比较倒
6	2	4.0	13.0	2.18	0.05	0.06	2.17	0.05	0.06	2. 15	0.06	0.06	2. 14	0.06	0.07	0	<b>Ø</b>	95	本発明
7	2	6.0	14.0	2.07	0.05	6.09	2.07	0.05	0.09	2.05	0.06	0.09	2.05	0.06	0.09	0	•	91	
8	2	6. D	13.0	2.15	0.05	0.06	2. 15	0.05	0.06	2. 13	0.05	0.05	2.12	0.06	0.07	0	6	91	
9	2	6.0	9.0	1.85	0.04	0.05	1.84	0.05	0.05	1.83	0.05	0.06	1.83	0.06	0.06	0	0	91	a
10	2	6.0	7. D	0. 14	8. CD	0.00	9. 14	0.00	0.90	0.14	0.00	6.40	0.14	0.00	0.00	0	×	91	比較例
11	2	7.0	13.0	2. 14	0.06	0.07	2. 12	0.06	0. 07	2. 13	0.06	0.08	2. 10	0. 07	8. 98	Δ	0	89	本発明
12	2	8.0	13.0	1.95	0.08	0.15	1.91	0.10	9. 16	1.86	0.15	0.18	1_80	0. 18	0.21	×	0	73	比較例

[0125] In addition, since the blinding of a processing liquid supply means occurred on the 2nd day of the consecutive processing, consecutive processing was stopped by the art 5.

[0126] Sufficient concentration is obtained, fogging is also suppressed low and by supplying pH seven or less liquid and pH eight or more liquid to a sample, respectively, and mixing them directly from Table 1, shows that generating of a sludge can be prevented. Moreover, pH of the liquid containing a color development chief remedy is four or less, and it is a book.

[0127] Furthermore, stable consecutive processing can be performed, by making it the method with which both mind space, the shelf life of processing liquid also improves and by supplying at least one of the two of both strong solutions to sensitive material through space shows that it is desirable.

[0128] the 440nm spectrum of the maximum blue reflection density Dmax (Y) (440nm) and the unexposed section of the sample at the time of the consecutive-processing start which the amount of supply was changed as shown in Table 2, and processed it like the example 1 about example 2 art 2 — reflection density Dmin (Y) and a 660nm spectrum — reflection density Dmin (C) was measured A result is shown in Table 2.
[0129]

[Table 2]

			発色現	見像液	発色現像液の						
実験	処理	発色	- 20	DрН	供給	量(=1/	<b>∕</b> ∎²)	Dmax(Y)	Dain(Y)	Dmin(C)	備考
No.	方法	現像液	部分	部分	部分	部分	A41	DWAX(Y)	DEFICT	Davide	1487
			液A	液Β	液A	液B	合計				
13	2	2	1.5	13	25	5	30	2.05	0.04	0.04	本発明
14	2	2	1.5	13	25	25	50	2.27	0.05	0.06	B
15	2	2	1.5	13	25	100	125	2.16	0.05	0.06	ø
16	2	2	1.5	13	25	140	165	2.05	0.05	0.06	,,
17	2	2	1.5	13	25	160	185	2.01	0.06	0. 06	"
18	2	2	1.5	13	100	25	125	2.04	0.06	0. 06	v
19	2	2	1.5	13	100	100	200	2.25	0.06	0. 07	ע
20	2	2	1.5	13	100	140	240	2.08	0.06	0. 07	#
21	2	2	1.5	13	100	160	260	2.06	0.08	0. 09	
22	2	2	1.5	13	140	100	240	2. 20	0.06	0.08	ø
23	2	2	1.5	13	140	140	280	2.14	0.07	0.09	
24	2	2	1.5	13	140	160	300	2.09	0.09	0.10	a
25	2	2	1.5	13	160	100	260	2.22	0.10	0.10	it
26	2	2	1.5	13	160	140	300	2.21	0.10	0.10	v
27	2	2	1.5	13	160	160	320	2.16	0.10	0.10	u

[0130] By this, wh in the amount of supply of each liquid to sensitive material is 5 - 150 ml/m2, required concentration is obtained, and it turns out that fogging is suppressed low.

[0131] In xample 3 art 2, the concentration of the color development chief remedy of the partial liquid A of color development liquid -2 was changed as shown in Table 3, and the maximum blue reflection density Dmax (Y) (440nm)

and the part light reflex concentration Dmin of 660nm of the unexposed sction (C) of the sample at the time of the consecutive-processing start which set pH of partial liquid B to 13.0, and processed it like the example 1 w r measured. Moreover, it saved at -5 degrees C, enclosing partial liquid A with a processing container, and 3, 10, 20, and 30 days after, the container was observed and the following criteria estimated low-temperature deposit nature. A result is shown in Table 3.

[0132] O A sludge is x by which the sludge was checked ten days [ from which the sludge was checked 20 days / which will not b checked / after O: ] after \*\*: also after [ of : ] 30 days. : The sludge was check d three days after.

[0133] [Table 3]

実験	発色現像	液のpH	発色現像:	主楽の濃度	Dmax(Y)	Dmin(C)	低温析出性	備考
No.	部分液A	部分液B	9/8	mol/ £	Dillax(1)	Diffill(O)		165-2
28	1.5	13.0	2	0.0046	2.02	0.04	0	本発明
29	1.5	13.0	5	0.011	2.05	0.04	0	本発明
30	1.5	13.0	30	0.069	2.23	0.05	0	本発明
31	1.5	13.0	40	0.092	2.27	0.05	0	本発明
32	1.5	13.0	50	0.11	2.25	0.06	0	本発明
33	1.5	13.0	100	0.23	2.15	0.06	0	本発明
34	1.5	13.0	200	0.46	2.10	0.06	0	本発明
35	1.5	13.0	400	0.92	2.08	0.07	0	本発明
36	1.5	13.0	450	1.03	2.01	0.09	Δ	本発明

[0134] If pH of the partial liquid A of color development liquid -2 is made or less into seven and concentration of a color development chief remedy is carried out [ l. ] in 0.005-1.00 mols /by this, it turns out that the problem of a deposit does not occur even if required concentration is obtained, and fogging is small and being saved at low temperature.

[0135] It is made to change, as the color development chief remedy of example 4 color-development liquid -2, the concentration of potassium carbonate, and the amount of supply of each liquid are shown in Table 4. Except having adjusted pH of partial liquid A and partial liquid B to 1.5 and 13.0, respectively the 440nm spectrum of the maximum blue reflection density Dmax (Y) (440nm) and the unexposed section of the sample at the time of the consecutive-processing start processed like the example 1 -- reflection density Dmin (Y) and a 660nm spectrum -- reflection density Dmin (C) was measured

[0136] A result is shown in Table 4.

[0137]

[Table 4]

実験	主薬濃度	炭酸加沙仏の	処理液	共給量(mg	/m²)	D00	Design	D-i-(C)	備考	
Nο.	(g/l)	濃度(g/ℓ)	部分液A	部分液B	比率	Dmax(Y)	Dmin(Y)	Dmin(C)	1875	
37	400	400	2.5	2.5	1	2.02	0.06	0.05	本発明	
38	400	400	2.5	5	2	2.09	0.06	0.05	本発明	
39	400	400	2.5	25	10	2.18	0.06	0.05	本発明	
40	400	400	2.5	100	25	2.14	0.06	0.05	本発明	
41	400	400	2.5	250	100	2.05	0.08	0.07	本発明	
42	400	400	2.5	300	120	2.04	0.10	0.09	本発明	
43	40	90	25	25	1	2.27	0.05	0.06	本発明	
44	40	90	25	50	2	2.21	0.05	0.06	本発明	
45	40	90	25	250	10	2.12	0.07	0.08	本発明	
46	40	90	25	275	11	2.09	0.09	0.10	本発明	

[0138] Table 4 shows that required conc ntration can be obtained and fogging can be low suppr ssed if the ratio of

the amount of supply of low pH liquid and high pH liquid is in the range of 1–100 by capacity.

[0139] Except having changed the color development processing time, as shown in Table 5, and having adjusted pH of the partial liquid A and B of color development liquid –2 to 1.5 and 13.0 in example 5 art 2, respectively the 440nm spectrum of the maximum blue reflection density Dmax (Y) (440nm) and the unexposed section of the sample at the time of the consecutive-processing start processed like the sample 1 — reflection density Dmin (Y) and a 660nm spectrum — reflection density Dmin (C) was measured A result is shown in Table 5.

[0140]

[Table 5]

実験Na.	処理期(秒)	Dmax(Y)	Dmin(Y)	Dmin(C)	備考
47	4	1.98	0.04	0.04	本発明
48	5	2.08	0.04	0.05	本発明
49	10	2.27	0.05	0.06	本発明
50	30	2.28	0.05	0.06	本発明
51	40	2.27	0.07	0.07	本発明
52	50	2.27	0.10	0.11	本発明

[0141] Table 5 shows that required concentration can be obtained and fogging can be low suppressed if development time is made into the range for 5 - 45 seconds.

[0142] It is made to change in example 6 art 2, as the concentration of the potassium carbonate of the partial liquid B of color development liquid -2 is shown in Table 6. Except having adjusted pH of partial liquid A and partial liquid B to 1.5 and 13.0, respectively the 440nm spectrum of the maximum blue reflection density Dmax (Y) (440nm) and the unexposed section of the sample at the time of the consecutive-processing start processed like the example 1 — reflection density Dmin (Y) and a 660nm spectrum — reflection density Dmin (C) was measured Moreover, low-temperature deposit nature was evaluated like the example 3 about partial liquid B. A result is shown in Table 6. [0143]

[Table 6]

実験No.	炭酸加	沙鴻度	Dmax(Y)	Dmin(C)	低温析出性	備考	
关款140.	g/4	mol/ £	Dillax(1)	Dillingo		5 MIN	
53	10	0.072	2.04	0.05	0	本発明	
54	15	0.11	2.16	0.05	0	本発明	
55	50	0.36	2.23	0.06	0	本発明	
56	90	0.66	2.27	0.06	Ø	本発明	
57	160	1.16	2.24	0.06	0	本発明	
58	450	3.26	2.08	0.07	0	本発明	
59	500	3.62	2.01	0.09	Δ	本発明	

[0144] Table 6 shows that a good result can be obtained if concentration of the alkali chemicals of high pH liquid is made into I., 0.1–4.5 mols /, and high concentration.

[Effect of the Invention] By this invention, it excels in dry feeling at quick processability, and the shelf life of development liquid can also obtain the art of a good silver-halide photosensitive material.

[Translation done.]

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#### **DESCRIPTION OF DRAWINGS**

[Brief Description of the Drawings]

- [Drawing 1] The outline block diagram of the principal part of the auto-processor used in the example.
- [<u>Drawing 2]</u> The perspective diagram of the principal part of the auto-processor of <u>drawing 1</u> .
- [Drawing 3] The perspective diagram near the feed-hopper dryness prevention means of the auto-processor of drawing 1.
- [Drawing 4] The outline block diagram of the principal part of the auto-processor which has two development meanses.
- [Drawing 5] The schematic diagram of the processing tank part of the automatic processor for immersing development which can supply two kinds of processing liquid for the color developments.
- [Drawing 6] Another side is the schematic diagram of the color development section of the automatic processor with which one side immerses for it and supplies two kinds of processing liquid for the color developments through space.

[Drawing 7] The schematic diagram of the automatic processor which can be supplied to the emulsion side of sensitive material through space after mixing two kinds of processing liquid for the color developments. [Description of Notations]

- 10 Heating Means
- 11 Heating Drum
- 30 Second Heating Means
- 33 Heating Belt
- 50 Development Means
- 51 First Processing Liquid Container
- 52 Processing Liquid Supply Means
- 54 Circulating Pump
- 55 Filter
- 56 Second Processing Liquid Container
- 57 Rotator
- 70 Sensitive-Material Detection Means
- 80 Feed-Hopper Dryness Prevention Means

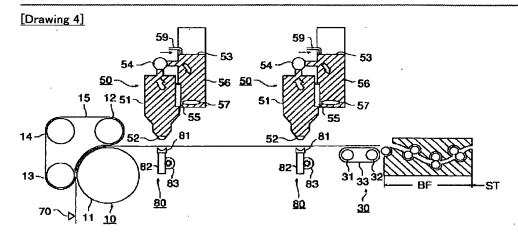
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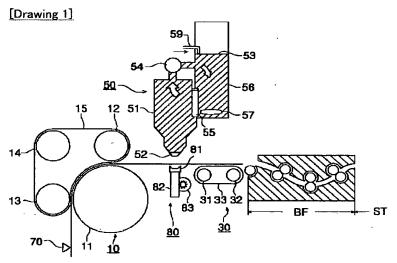
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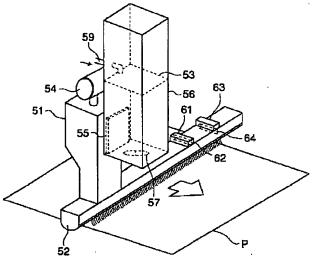
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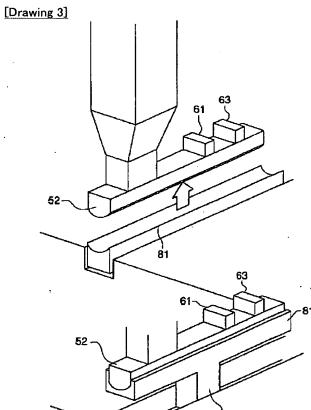
# **DRAWINGS**



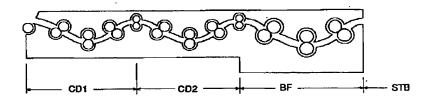


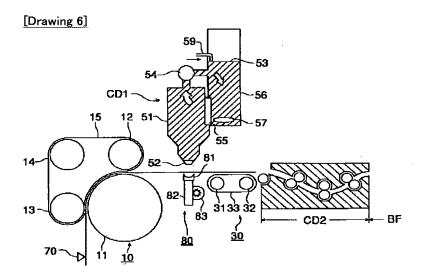
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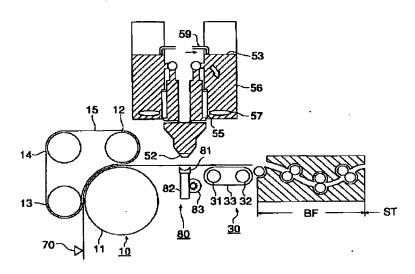


[Drawing 5]





# [Drawing 7]



[Translation done.]